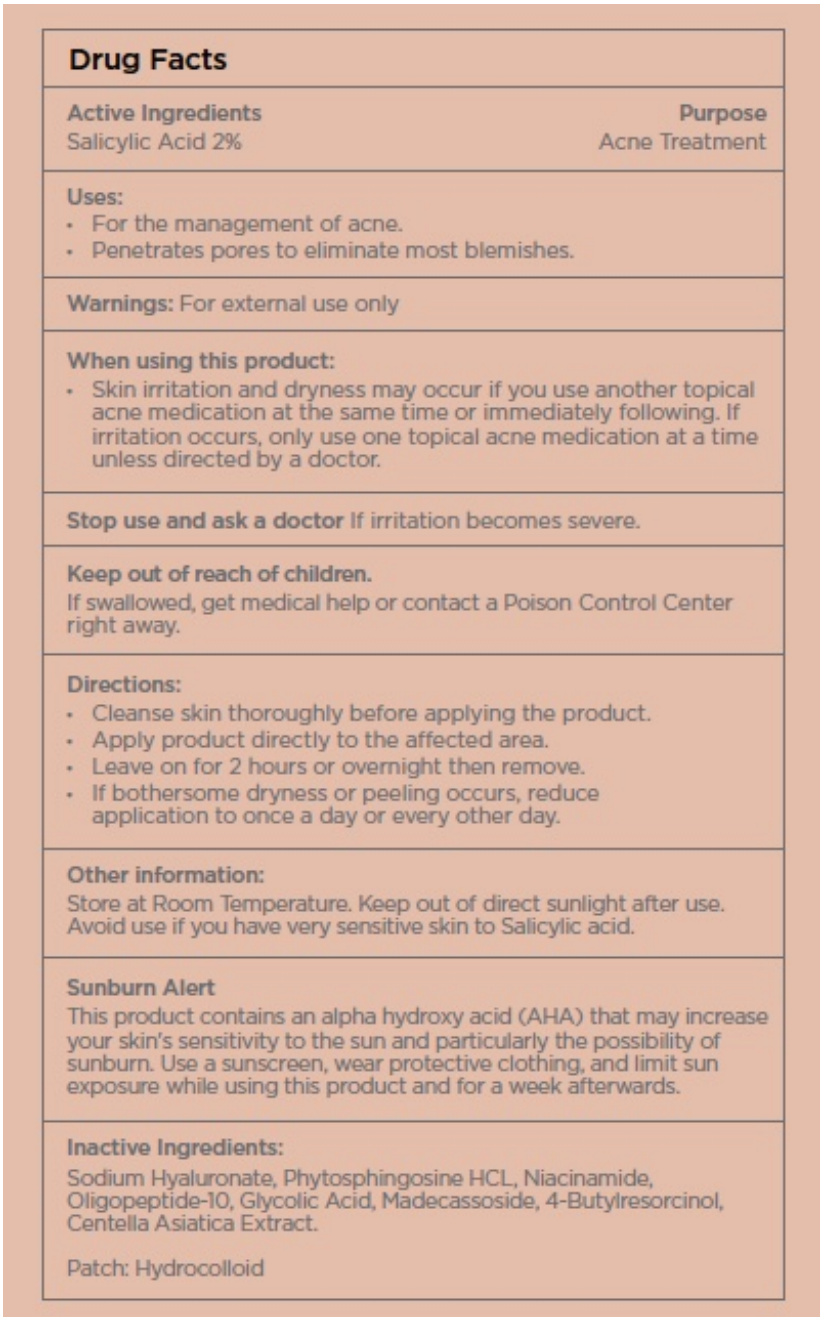
KILLA EXTRA STRENGTH- salicylic acid patch
Zitsticka, Inc.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Active Ingredients



Drug Facts

Active Ingredients — **Purpose**
Salicylic Acid 2% — Acne Treatment

Uses:
- For the management of acne.
- Penetrates pores to eliminate most blemishes.

Warnings: For external use only

When using this product:
- Skin irritation and dryness may occur if you use another topical acne medication at the same time or immediately following. If irritation occurs, only use one topical acne medication at a time unless directed by a doctor.

Stop use and ask a doctor If irritation becomes severe.

Keep out of reach of children.
If swallowed, get medical help or contact a Poison Control Center right away.

Directions:
- Cleanse skin thoroughly before applying the product.
- Apply product directly to the affected area.
- Leave on for 2 hours or overnight then remove.
- If bothersome dryness or peeling occurs, reduce application to once a day or every other day.

Other information:
Store at Room Temperature. Keep out of direct sunlight after use. Avoid use if you have very sensitive skin to Salicylic acid.

Sunburn Alert
This product contains an alpha hydroxy acid (AHA) that may increase your skin's sensitivity to the sun and particularly the possibility of sunburn. Use a sunscreen, wear protective clothing, and limit sun exposure while using this product and for a week afterwards.

Inactive Ingredients:
Sodium Hyaluronate, Phytosphingosine HCL, Niacinamide, Oligopeptide-10, Glycolic Acid, Madecassoside, 4-Butylresorcinol, Centella Asiatica Extract.

Patch: Hydrocolloid

Purpose

Acne Treatment

Ask a doctor

If irritation becomes severe

When using this product

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Killa Extra Strength PDP

KILLA EXTRA STRENGTH

salicylic acid patch

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:81746-418
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
	Basis of	

Ingredient Name	Basis of Strength	Strength
MAGNESIUM SALICYLATE (UNII: 41728CY7UX) (SALICYLIC ACID - UNII:O414PZ4LPZ)	MAGNESIUM SALICYLATE	0.0664 g in 3.32 g

Inactive Ingredients

Ingredient Name	Strength
OLIGOPEPTIDE-10 (UNII: Q46328TRNK)	0.1593 g in 3.32 g
NIACINAMIDE (UNII: 25X51I8RD4)	0.1593 g in 3.32 g
PHYTOSPHINGOSINE HYDROCHLORIDE (UNII: TT871XV7TU)	0.166 g in 3.32 g
GLYCOLIC ACID (UNII: 0WT12SX38S)	0.0232 g in 3.32 g
4-BUTYLRESORCINOL (UNII: 2IK4UQ3ZGA)	0.00664 g in 3.32 g
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	2.715 g in 3.32 g
MADECASSOSIDE (UNII: CQ2F5O6YIY)	0.0232 g in 3.32 g

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:81746-418-01	3.32 g in 1 BOX; Type 2: Prefilled Drug Delivery Device/System (syringe, patch, etc.)	01/01/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	M006	01/01/2023	

Labeler - Zitsticka, Inc. (117778611)

Establishment

Name	Address	ID/FEI	Business Operations
Raphas		695914964	manufacture(81746-418)

Revised: 6/2023 Zitsticka, Inc.